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(54) Title: PROCESS FOR INCREASING STAIN-RESISTANCE OF CATIONIC-DYEABLE MODIFIED POLYAMIDE FIBERS (57) Abstract This invention relates to an improved process for dyeing cationic-dyeable polyamide fibers, especially carpet fibers, with an acid dye to provide stain-resistant fibers. The polyamide fibers contain cationically dyeable sulfonate groups along the polymer chain. The improvement involves adding certain water-soluble salts to a dyebath at a concentration of at least 20 % based on the weight of fiber.		

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TITLE

Process for Increasing Stain-Resistance
of Cationic-Dyeable Modified Polyamide Fibers

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BACKGROUND OF THE INVENTIONField of the Invention

This invention relates to an improved process
for preparing stain-resistant polyamide fibers,
10 especially carpet fibers, containing cationically-
dyeable sulfonate groups along the polyamide polymer
chain, wherein the fibers have been dyed with an acid
dyestuff.

15 Description of the Related Art

As known in the art, polyamide fibers can be
made to resist staining by acid dyes by copolymerizing
in the polyamide polymer a small percentage of a
cationically dyeable monomer such as an aromatic
20 sulfonate, for example, 1 to 4 weight percent of the
sodium salt of 5-sulfoisophthalic acid. In Ucci, U.S.
Patent 4,579,762 nylon 6 and nylon 6,6 carpet fibers
formed from polymers having aromatic sulfonate units in
the polymer chain and having improved stain-resistance
25 to acid dyes are disclosed.

From Windley, U.S. Patents 5,164,261 and
5,468,554, it is further known that such cationic-
dyeable modified polyamide fibers can be made more
30 stain-resistant upon being dyed with at least 0.0048
weight percent of an acid dyestuff. However, under
some circumstances, especially when it is desirable to
dye such fibers to only a light shade, it has been
found that maximum stain-resistance (i.e., no staining)
35 may not be obtained.

Chao, U.S. Patent 5,030,246 discloses a
process for continuously dyeing polyamide fibers which
do not contain cationic-dyeable monomeric units in the
40 polymer chain. Rather, the fibers are coated with a

stainblocking agent which may contain aromatic sulfonate groups. The fibers can be acid dyed to a deeper shade by adding certain ammonium and metal salts to the dye liquor (typically in an amount from 0.2 to 5 8% based on the weight of the fiber). Chao discloses that effective salts include salts of lithium, calcium, and magnesium, as well as certain ammonium, sodium and potassium salts. The use of the salts results in greater uptake of the dyestuff from the dye liquor onto 10 the fibers, and the excess dye not taken up by the fibers is subsequently rinsed off with water.

In Jenkins, U.S. Patent 5,466,527, cationic dyeable nylon fibers containing a sufficient amount of 15 SO_3H groups or COOH groups within the polymer structure to render the nylon fiber dyeable with cationic dyes are disclosed. A process for improving the stain-resistance, lightfastness, and ozone-resistance of such fibers is disclosed, wherein the fibers are dyed with 20 level acid dyes or premetallized acid dyes at a pH value less than 7.0. In Example 6 (columns 13-14), a carpet made with cationic dyeable nylon fiber is described as being dyed in a dyebath containing level acid dyes or premetallized acid dyes and 2% sodium 25 sulfate (Glauber salt) based on weight of fiber. Tables I and II in Example 6 show that an improvement in dye exhaustion levels is obtained when 2% sodium sulfate is added to the dyebath over dye exhaustion levels obtained from dyebaths which do not contain 30 sodium sulfate. Thus, it is known from Jenkins that sodium sulfate can be added to a dyebath and better exhaustion of the acid dye onto the fiber can be obtained. However, it would be desirable to have a process for acid dyeing cationic-dyeable modified 35 polyamide fibers, wherein the stain-resistance of the fibers can be improved.

The present invention provides such a process. Particularly, it has now been discovered that when certain salts are included in the acid dyebath at certain concentrations, the stain-resistance of the acid dyed fibers is improved. The stain-resistance of the acid dyed fibers is improved without having to increase the dye content on the fiber.

10

SUMMARY OF THE INVENTION

The present invention provides an improved process for dyeing polyamide fibers with an acid dyestuff in a dyebath, wherein the fibers comprise a polyamide copolymer which contains cationic-dyeable aromatic sulfonate salt group substituents along the polymer chain. The improvement comprises adding a water-soluble salt selected from the group consisting of ammonium, potassium, and sodium salts, and mixtures thereof to a dyebath at a concentration of at least 20% based on the weight of the fiber to produce acid dyed fibers having a stain-resistance rating of at least 8.0 on the AATCC Red 40 Scale.

Preferably, the salt is selected from the group consisting of sodium and potassium chlorides; sodium, potassium, and ammonium sulfates; and sodium acetate. In some instances, the concentration of salt in the dyebath may be at least 100% based on weight of fiber. The acid dye may be a premetallized acid dye. The fibers may be acid dyed at a pH level greater than 6.5, although such a pH level is not necessary. Suitable polyamide copolymers for use in this invention include copolymers containing 1 to 4 weight percent of monomer units derived from a salt or other derivative of 5-sulfoisophthalic acid based on the final weight of the copolymer. The polyamide copolymer may also contain units selected from the group consisting of

polyhexamethylene adipamide units, poly-ε-caprolactam units, and mixtures thereof.

5 DETAILED DESCRIPTION OF THE INVENTION

As known in the art, polyamide fibers can be visibly and permanently stained when left in contact with a solution containing acid dyes typically used as colorants for foodstuffs. Those skilled in the art
10 have developed different methods for rendering polyamide fibers stain-resistant. For example, in the foregoing Windley, U.S. Patents 5,164,261 and 5,468,554, the entire disclosures of which are hereby incorporated by reference, the polyamide fibers are
15 rendered stain-resistant by copolymerizing in the polyamide polymer a cationic dye modifier and by adding an acid dye to the polymer melt or by dyeing the fibers with an acid dye from a dyebath. The present invention is an improvement over the process disclosed in the
20 foregoing patents and involves adding certain salts to the dyebath which selectively improve the stain-resistance of the dyed fibers without the need for increasing the dyestuff content on the fiber.

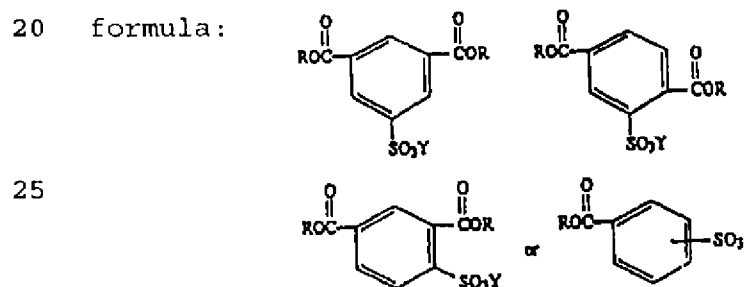
25 More particularly, this invention provides an improved process for dyeing polyamide copolymer fibers containing a cationic dye modifier with an acid dye, producing dyed fibers which, when dried, have a stain-resistance rating of at least 8.0, preferably 9.0, on
30 the AATCC Red 40 Stain Scale. The term, "acid dye" includes premetallized acid dyes. The improved process is accomplished by dyeing the cationic dyeable fibers with an acid dye in a dyebath which contains salt at a concentration to provide at least 20% salt based on the
35 weight of fibers (% owf) being dyed. The process is particularly advantageous for fabrics which are dyed to light or medium shades. It is presumed that the improvement is equally effective for fibers dyed to

deep shades, but stains are less evident on deep-dyed fibers, so the improvement is difficult to evaluate. The term "fiber" includes both continuous (e.g., bulked continuous filament) and short lengths (e.g., staple).

- 5 Effective salts are ammonium, sodium and potassium salts, or combinations of any of these salts, at a concentration of at least 20% based on weight of fiber. When these salts are used in accordance with this invention, dyeing with acid dyes takes place rapidly
 10 and there is no need to adjust the pH of the dyebath at any stage of the dying process. These acid-dyed cationically-modified polyamide fibers have greater stain-resistance than cationically-modified polyamide fibers which have been dyed in a dyebath which does not
 15 contain these salts.

Cationic Dyeable Modified Polyamides

The cationic dye modifier used to form the polyamide copolymers useful in this invention have the
 20 formula:



- where Y is H, Li, Na, K, or Cs and R is H or an alkyl group containing 1 to 5 carbon atoms. The -OR groups are lost during polymerization. The preferred cationic dye modifiers are those containing two carboxyl groups with 5-sulfoisophthalic acid being especially preferred. Generally, sufficient cationic dye modifier
 35 is used to produce a copolymer containing from 1 to 4 weight percent (wt.%) of cationic dye modifier based on the final polymer weight of the polyamide copolymer with from 2 to 3 wt.% being the preferred range.

- 40 The polyamide copolymers useful in this invention can be prepared by blending the salt of the

base polyamide (e.g., nylon 6,6 salt or caprolactam) with the cationic dye modifier followed by polymerizing the blended composition in an autoclave using standard polymerization procedures, solidifying and fragmenting the polymer, increasing the degree of polymerization by further polymerizing the polymer in the solid phase while simultaneously super-drying the polymer, and further polymerizing the polymer in the molten state. In a preferred embodiment used for production of carpet fibers, the copolymer is prepared by polymerizing nylon 6,6 salt containing sulfonate modifier in an autoclave to a relative viscosity of about 35, increasing the relative viscosity to about 45 to 65 by solid-state polymerization while super-drying the polymer, and melting the polymer in a screw extruder, transfer line and spin block to produce a copolymer with a relative viscosity between 45 and 70. In the production of textile fibers, the acceptable relative viscosity can be lower (about 35 to 45). Preferably, the copolymers used in this invention contain from 20 to 40 amine ends per 10^6 grams of copolyamide.

Dyeing Procedure

A dyebath is prepared by adding the desired amount of acid dye and salt to water. Dye concentrations can vary from 0.01 to 0.05%, based on weight of fiber. The salt concentration must be sufficiently high to exhaust the dyestuff from the dyebath onto the fiber. As stated earlier, it has been found that this salt concentration must be at least 20%, based on weight of fiber (% owf), and can be as high as 200%, or even higher, for the salts which are useful in this invention. The fibers may be dyed at a pH level greater than 6.5, but such a pH level is not necessary. If desired, a lower pH level can be used.

The results reported in the following examples were obtained by placing 10 grams of fiber in 200 ml of

dyebath containing 0.0037 grams of dye and salt at various concentrations. The dyebath was heated to boiling in the shortest possible period and held at the boil for 5-30 minutes, usually about 10 minutes. The
5 dyebath was then cooled to 70-80°C. Cold water may be added to accomplish the cooling. The fibers were removed, washed and dried, either at room temperatures or by heating. The fibers were dyed to a light gray shade. Comparable results were obtained with fibers
10 dyed to dune and beige shades. Equivalent results were obtained for fibers in the form of yarns, or woven, knitted, or pile fabrics. This procedure can be readily adapted to a continuous process.

15 The following examples are illustrative of this invention but should not be construed as limiting the scope of the invention.

TEST METHODS

20 Stain Test Method

The following test procedure was used to determine the stain-resist performance of the fiber samples.

25 A solution of staining agent was prepared by dissolving 45 grams of cherry-flavored, sugar-sweetened "Kool-Aid" premix powder in 500 cc. of water. The solution allowed to reach room temperature, i.e., 75°F ± 5°F (24°C ± 3°), before using. The colorant used in
30 the "Kool Aid" solution was Red Acid Dye 40.

A specimen, approximately 1.5 x 3 inches (3.8 x 7.6 cm.) for carpet samples and approximately 2 x 4 inches (5.1 x 10.2 cm.) for knitted fabrics was cut
35 from each sample and placed on a flat non-absorbent surface. The solution of staining agent was poured onto each specimen through a cylinder to form a 1 to 2 inch (2.5 to 5.1 cm.) circular stain, using 20 cc. for

carpet samples and 10 cc. for textile samples. The samples were lagged in the laboratory for twenty-four hours and then rinsed thoroughly with cool tap water and squeezed dry, using an extractor to remove excess
5 solution.

The stain-resistance of the specimen was determined visually according to the color left on the stained area of the sample. The color depth was
10 determined by comparison with a series of ten transparent plastic rectangles in accordance with the AATCC Red 40 Stain Scale, in which 10 represents no staining, 9 very light staining, with increasing color as the scale decreases to 1, which represents heavy
15 staining.

In each of the following examples, 200 ml. of dyebath and 10 grams of fiber sample were used. Concentrations of salt and dye are expressed as
20 percentages based on the weight of fiber (% owf). The same weight of dye (0.0037 gram) was used in each example. In all but one of the examples, the cationic dyeable polyamide copolymer contained 3% by weight of 5-sulfoisophthalate. The dyed samples were light gray
25 in color. The dye formulation was composed of the following ingredients: 0.015% owf Tectilon yellow 3R KWL 200 (acid yellow 246), 0.0075% owf Tectilon red 2B KWL 200 (acid red 361), and 0.005% owf Tectilon blue 4RS KWL 200. The pH levels of the solutions were
30 measured with a Fisher Accumet pH Meter Model 610A equipped with a Fisher glass electrode.

EXAMPLES

Fiber Preparation

35 A polyamide copolymer was prepared by blending nylon 6,6 salt and the sodium salt of 5-sulfoisophthalic acid and polymerizing in an autoclave. The polymer melt was solidified, fragmented, and

polymerized further in the solid state in an inert atmosphere at a temperature of 185°C. The copolymer was then fed to a twin screw extruder and discharged into a transfer line at a temperature of 290°C. It was
5 extruded through a spinneret to produce yarns in which each of 128 filaments had four symmetrically-placed voids. After application of finish, the yarns were drawn 2.7X at 190°C in a continuous process. The drawn filaments were passed through a jet where they were
10 impinged with air at 240°C and 120°C and collected on a screen drum. The yarn was removed by a take-up roll and wound onto tubes. The knitted and tufted fabrics used in the following Examples were prepared from these yarns.

15

Example 1

Certain salts, such as calcium salts, are effective in exhausting the dyestuff from the dyebath onto the fiber at low and high salt concentrations.
20 However, the stain-resistance of fiber samples dyed to light and medium shades by methods using such salts is unacceptable, as shown in Table 1. In comparison, the process of this invention employs certain salts at a concentration of at least 20% on weight of fiber (%
25 owf) and provides fibers having good stain-resistance, as shown in Table 1.

TABLE 1

Salt		pH of Soln	% Dyestuff on Fiber	Stain Rating
Composition	% OWF			
*Ca(H ₂ PO ₄) ₂	2	5.3	0.037	6.0
*Ca(H ₂ PO ₄) ₂	20	5.3	0.037	6.0
* (NH ₄) ₂ SO ₄	10	7.2	Less than 0.037	7.0
*Na ₂ SO ₄	10	7.8	Less than 0.037	6.0
Na ₂ SO ₄	20	7.4	Less than 0.037	8.0
(NH ₄) ₂ SO ₄	40	6.8	0.037	8.0

*Comparative Examples

5 Example 2

When the salts of this invention are used in the dyebath at concentrations above 20%, based on weight of fiber (% OWF), they are effective in imparting even greater stain-resistance to cationic-dyeable modified polyamide fibers which are dyed with an acid dyestuff. Such fibers are made from polyamide copolymer containing 2% and 3% 5-sulfoisophthalate. This effect is illustrated in Table 2 for polyamide fibers taken from knitted fabrics which have been dyed to a light gray color, wherein the fibers are made from polyamide copolymer containing 2% and 3% 5-sulfoisophthalate.

TABLE 2

% Sulfoisophthalate	Salt		pH	% Dyestuff on Fiber	Stain Rating
	Composition	% OWF			
2	NaCl	185	7.4	0.037	9.0
3	KCl	185	7.4	0.037	9.0

20 Example 3

The process of this invention is useful for dyeing and imparting good stain-resistance to the above-described polyamide fibers in any form e.g., yarns or nonwoven, knitted, woven or pile fabrics.

This effect is illustrated in Table 3 for fibers dyed to a light gray shade.

TABLE 3

Substrate	Salt		pH	% Dyestuff on Fiber	Stain Rating
	Composition	% OWF			
Yarn	NaCl	200	7.4	0.037	9.0
Knitted Fabric	NaCl	185	7.4	0.037	9.0
Pile Fabric (carpet)	$(\text{NH}_4)_2\text{SO}_4$	40	6.8	0.037	8.0

5

Example 4

The data in Table 4 illustrate the unique effectiveness of the sodium and potassium salts for producing polyamide fibers which have good stain-resistance after being dyed with the above-described dyes, as compared to similar salts of lithium, magnesium, zinc, and calcium. All of the polyamide fibers were made from copolymers containing 3% by weight of 5-sulfoisophthalic acid and were dyed to a light gray shade. The tests were run on fabric taken from knitted socks.

10

15

TABLE 4

Salt		pH	% Dyestuff on Fiber	Stain Rating
Composition	% OWF			
Na_2SO_4	200	7.4	0.037	9.0
K_2SO_4	200	7.9	0.037	9.0
* Li_2SO_4	200	8.3	0.037	6.0
* MgSO_4	200	7.3	0.037	6.0
* ZnSO_4	200	6.8	0.037	6.0
* $\text{Ca}(\text{H}_2\text{PO}_4)_2$	20	5.3	0.037	6.0
Na_2SO_4	185	7.8	0.037	9.0
NaOAc	110	7.8	0.037	9.0
KCl	185	7.4	0.037	9.0

* Comparative Examples

CLAIMS:

1. An improved process for dyeing polyamide fibers with an acid dyestuff in a dyebath, wherein the fibers comprise a polyamide copolymer which contains cationic-dyeable aromatic sulfonate salt group substituents along the polymer chain, and wherein the improvement comprises adding a water-soluble salt selected from the group consisting of ammonium, potassium, and sodium salts, and mixtures thereof to the dyebath at a concentration of at least 20% based on the weight of the fiber to produce acid dyed fibers having a stain-resistance rating of at least 8.0 on the AATCC Red 40 Stain Scale.
2. The process of claim 1, wherein the salt is selected from the group consisting of sodium and potassium chlorides; sodium, potassium, and ammonium sulfates; and sodium acetate.
3. The process of claim 2, wherein the concentration of salt in the dyebath is at least 100% based on the weight of the fibers.
4. The process of claim 1, wherein the acid dye is a premetallized acid dye.
5. The process of claim 1, wherein the fibers are dyed at a pH level greater than 6.5.
6. The process of claim 1, wherein the polyamide copolymer contains 1 to 4 weight percent of monomer units derived from a salt, or other derivative of 5-sulfoisophthalic acid based on the final weight of the copolymer.

7. The process of claim 1, wherein the polyamide copolymer contains units selected from the group consisting of polyhexamethylene adipamide units, poly- ϵ -caprolactam units, and mixtures thereof.

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INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 D06P3/24 D06P1/673 D06P1/653

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 D06P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 92 02570 A (DU PONT) 20 February 1992 see page 5, line 3 - page 6, line 5; claims & US 5 164 261 A (WINDLEY) cited in the application ---	1-7
A	DATABASE WPI Section Ch, Week 8948 Derwent Publications Ltd., London, GB; Class A23, AN 89-350923 XP002042578 & JP 01 260 061 A (TEIJIN LTD) , 17 October 1989 see abstract --- -/--	1-7

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/08409

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 013, no. 546 (C-661), 6 December 1989 & JP 01 223908 A (TEIJIN LTD), 7 September 1989, see abstract ---	1-7
A	WO 91 17301 A (BURLINGTON INDUSTRIES INC) 14 November 1991 see claims & US 5 466 527 A (JENKINS) cited in the application ---	1-7
A	US 5 030 246 A (CHAO NELSON P C) 9 July 1991 cited in the application see column 2, line 61 - line 68; claims; table 1 ---	1-5
A	GB 2 160 230 A (SNIA BPD SPA; SNIA FIBRE) 18 December 1985 see page 4, line 29 - line 55; claims ---	1-7
A	FR 2 325 761 A (SANDOZ SA) 22 April 1977 see claims -----	1-5

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/08409

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9202570 A	20-02-92	US 5164261 A AU 8444491 A US 5468554 A	17-11-92 02-03-92 21-11-95
WO 9117301 A	14-11-91	US 5085667 A AU 6909891 A DE 69023514 D DE 69023514 T EP 0527129 A US 5350426 A US 5354342 A US 5466527 A US 5571290 A US 5199958 A	04-02-92 27-11-91 14-12-95 18-07-96 17-02-93 27-09-94 11-10-94 14-11-95 05-11-96 06-04-93
US 5030246 A	09-07-91	NONE	
GB 2160230 A	18-12-85	BE 902635 A DE 3520922 A NL 8501640 A	11-12-85 12-12-85 02-01-86
FR 2325761 A	22-04-77	CH 610175 A CA 1095202 A DE 2641159 A GB 1538963 A JP 1227391 C JP 52040682 A JP 58045517 B US 4181498 A BE 846504 A	12-04-79 10-02-81 31-03-77 24-01-79 31-08-84 29-03-77 11-10-83 01-01-80 23-03-77